Ansell Perry

MAR 10 2000

Next Step Powder Free Latex Examination Gloves (Protein Label Claim)
Ansell Perry
1875 Harsh Avenue SE
Massillon, Ohio 44644
Telephone: 330-833-2811
Fax: 330-833-6213

Checklist
Section 21.0

K000165

[1] 510 (k) Summary [Revised]

[2] Ansell Perry Inc.
1875 Harsh Avenue SE
Massillon, Ohio 44646

Telephone: 330-833-2811
Fax: 330-833-6213

Contact: James R. Chatterton
Telephone: 330-833-2811
Fax: 330-833-6213

January 19, 2000

[3] Trade Name: Next Step Powder Free Latex Examination Gloves (Protein Label Claim)
Common Name: Examination Gloves
Classification Name: Patient Examination Glove

[4] Next Step Powder Free Latex Examination Gloves (Protein Label Claim), meet all of the requirements of ASTM D 3578-99.

[5] Next Step Powder Free Latex Examination Gloves (Protein Label Claim) meet all the current specifications for ASTM D 3578-99 Rubber Examination Gloves.

[6] Next Step Powder Free Latex Examination Gloves (Protein Label Claim) are disposable device intended for medical purposes that is worn on the examiners hand to prevent contamination between patient and examiner.

[7] Next Step Powder Free Latex Examination Gloves (Protein Label Claim) are summarized with the following technological characteristics compared to ASTM or equivalent standards.

Characteristics	Standard
Dimensions	Meets ASTM D 3578-99
Physical Properties	Meets ASTM D 3578-99

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Freedom from holes	Meets ASTM D 3578-99 Meets ASTM D 5151-99
Powder-Free Meets described test in ASTM D 6124-97	Not more than 2 mg residue by mass.
Protein Label Claim	This latex glove contains 50 micrograms or less of total water extractable protein per gram.
Biocompatability	
Primary Skin Irritation in Rabbits	Passes
Guinea Pig Sensitization	Passes

[8] The performance test data of the non clinical tests are the same as mentioned immediately above.

[9] Clinical data is not needed for medical gloves or for most devices cleared by the 510(k) process.

[10] It is concluded that Next Step Powder Free Latex Examination Gloves (Protein Label Claim) are as safe, as effective, and perform as well as the glove performance standards referenced in Section 7 above and therefore meet:

ASTM listed standards,
FDA hole requirements, and
labeling claims for the product.

[11] This summary will include any other information reasonably deemed necessary by The FDA.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAR 10 2000

Mr. James R. Chatterton
Vice President Regulatory
Ansell Perry, Inc.
1875 Harsh Avenue, S.E.
Massillon, Ohio 44646

Re: K000165
Trade Name: Next Step Powder Free Latex Examination Gloves (contains 50 mcgm or less of total extractable protein per gram)
Regulatory Class: I
Product Code: LYY
Dated: February 24, 2000
Received: February 25, 2000

Dear Mr. Chatterton:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Good Manufacturing Practice for Medical Devices: General (GMP) regulation (21 CFR Part 820) and that, through periodic GMP inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of

the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4690. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "http://www.fda.gov/cdrh/dsmamain.html".

Sincerely yours,

Timothy A. Ulatowski
Director
Division of Dental, Infection Control,
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

3.0 Indications for Use Statement:

INDICATIONS FOR USE

Applicant: Ansell Perry Inc.

510(K) Number (if known): K000165 *
(Next Step)

Device Name: Patient Examination Glove, Powder Free with Protein Label Claim LATEX
Contains 50 mcgm or less of Total Water Extractable Protein per gram

Indications For Use:

A disposable device intended for medical purpose that is worn on the examiners hand to prevent contamination between patient and examiner.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH Office of Device Evaluation (ODE)

Olin S. Lin

(Division Sign-Off)
Division of Dental, Infection Control,
General Hospital Devices
510(k) Number K000165

Prescription Use _____ OR Over-The-Counter X
21 CFR 801.109

(Optional Format 1-2-96)